



**Group 1 : rocks**

1. Andesite
2. Ophiolite
3. Gabbro
4. Peridotite

**Group 2 : characteristics**

- a. Magmatic rock with grain structure belonging to oceanic crust.
- b. Rock complex belonging to oceanic lithosphere.
- c. Magmatic rock with a microlite structure characterizing subduction zones.
- d. Magmatic rock with a grain structure belonging to an upper mantle.
- e. Magmatic rock with a grain structure characterizing collision zones.

**Section II: Scientific reasoning and communication in graphic and written modes (15 pts)**

**Exercise 1 (3, 25 pts)**

During a short and intense exercise, such as in sprinting, the muscle power developed is very important. Therefore, ATP regeneration depends on a number of metabolic reactions. In order to understand the relationship between these reactions and how the body provides a sprinter with energy, we suggest the following data:

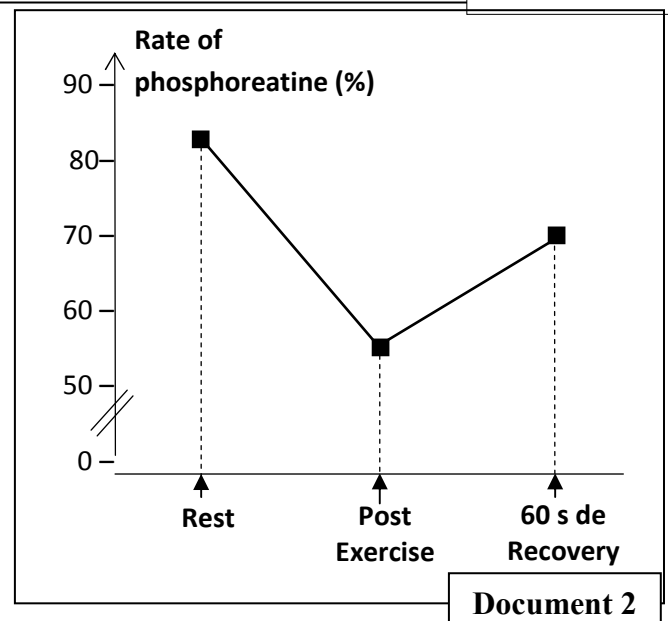
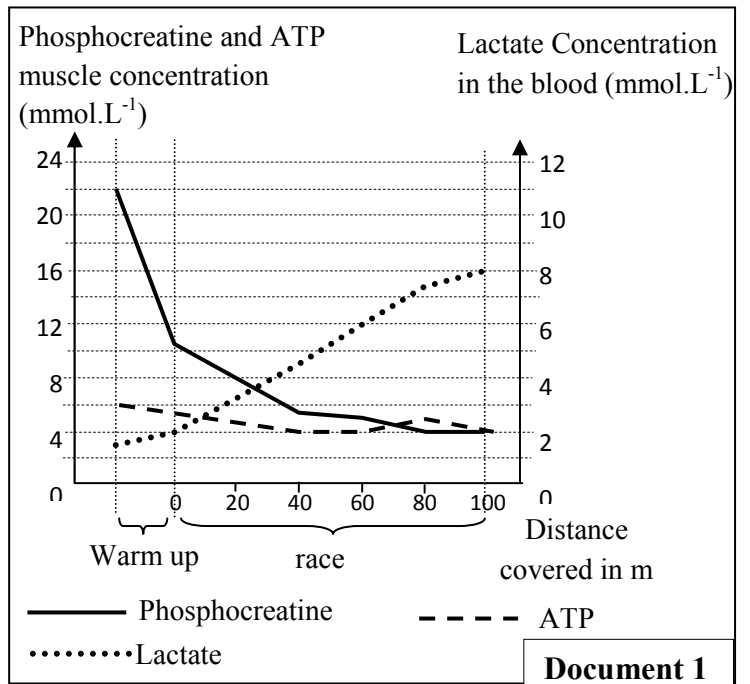
- During a warm up and a 10-second race, we have measured the variation of ATP concentration, phosphocreatine at the muscle level and blood lactate concentration in the sprinter. The results are presented in document 1.

1. **Describe** the concentration variation of ATP, lactate and phosphocreatine in the sprinter (document 1), then **give** an explanation of the ATP origin during this physical exercise. (1pt)

- We have measured the evolution of phosphocreatine levels in a sportsman's muscular biopsies collected while resting, after 45-second exercise leading to exhaustion (post exercise), and after 60-second recovery. Document 2 presents the obtained results.

2. Bearing in mind that recovery is achieved thanks to the contribution of Oxygen carried by blood, **suggest a hypothesis** to explain phosphocreatine evolution after the 60-second recovery. (0.25 pts)

- Document 3 presents the follow-up results of the concentration of the three phosphate compounds in a sportsman (PCr, ATP and inorganic phosphate « Pi ») before physical exercise, during a short-term physical exercise and after recovery. Document 4 explains the relation between ATP and phosphocreatine.

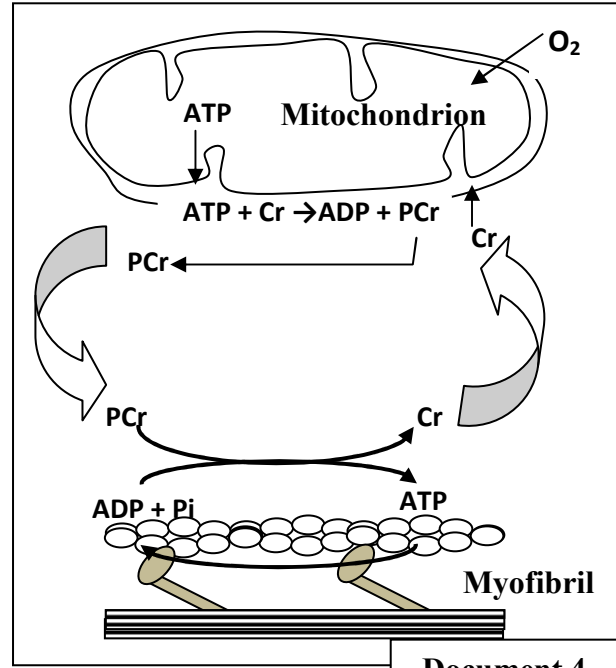


	before exercise	during exercise	after recovery
Pi	+	+++	+
ATP	++	++	++
PC	+++	++	+++

+ : low concentration ; ++ : average concentration ;  
+++ : high concentration

**Document 3**

- What pieces of information can be extracted from document 3. (0.75 pts)
- Based on your answer to question 3 and by exploiting the data in document 4, show the relationship between the variations of the three compounds in the sprinter during exercise and after recovery, and then verify the formulated hypothesis (answer to question 2). (1,25pt)



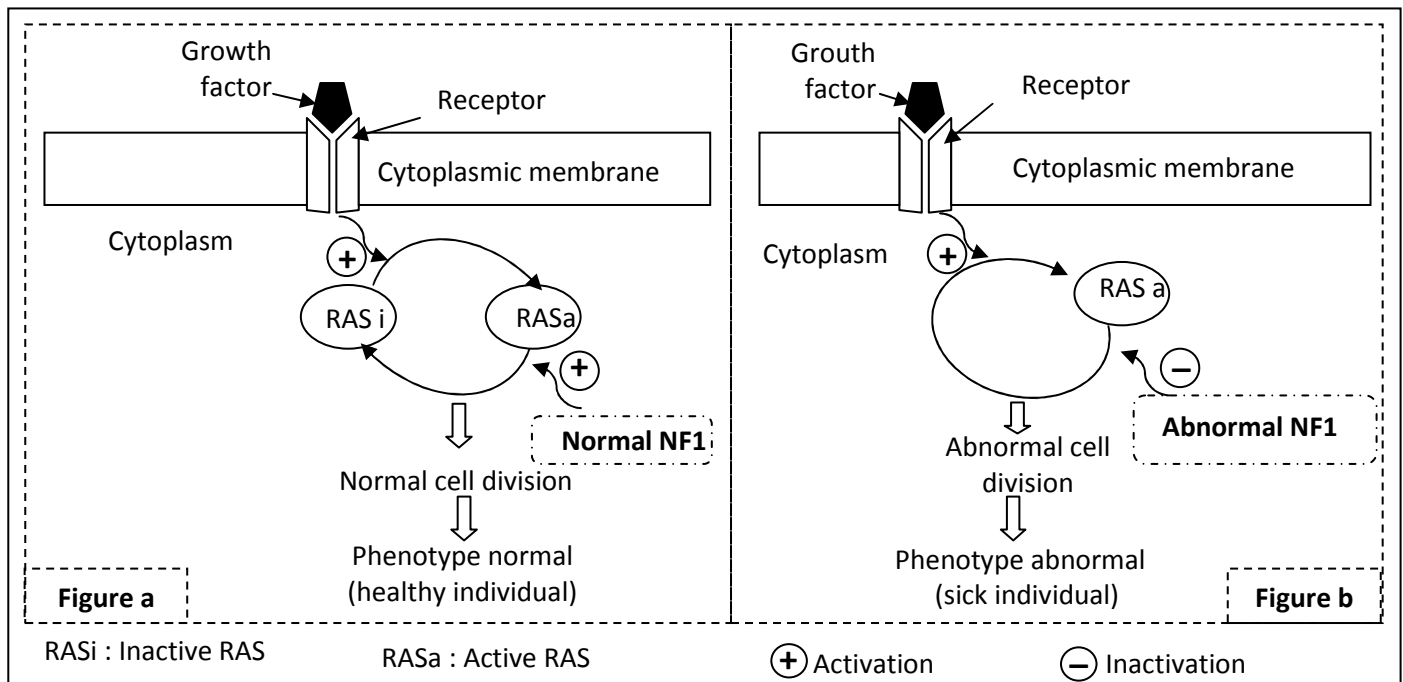
**Document 4**

**Exercise 2 (4,75 pts)**

Neurofibromatosis type 1 is a hereditary disease. Some symptoms of this disease include: Pigmented spots on the skin (café-au-lait spots) with overall risk of benign tumor, neurofibroma, and poor skeletal formation.

To determine the genetic origin of this disease, we suggest the following data:

❖ Neurofibromatosis type 1 is associated with a protein called Neurofibromin 1 (NF1). This protein controls the activity of another protein called **RAS**, which interferes in cell division and multiplication. Protein NF1 is found in two forms, one normal and the other abnormal. The two figures **a** and **b** in document 1 represent the relation between protein NF1, RAS protein activity and the nature of cell division in a healthy person (figure a) and in a sick person (figure b).



**Document 1**

1. Use document 1 to **compare** the effect of NF1 protein on RAS protein in the healthy person and in the sick person, then **show** the protein-trait relationship. (1 pt)

❖ Protein NF1 synthesis is controlled by a gene called (NF1) which exists in two allelic forms. Document 2 presents a transcribed strand fragment of a normal allele in a healthy person and a transcribed strand fragment of an abnormal allele in a sick person. Document 3 presents an extract of the genetic code table.

Number of triplet	6531	6532	6533	6534	6535	6536
Fragment of a normal allele	AAA	ACG	AAA	CTG	TAG	GAA
Fragment of an abnormal allele	AAA	ACG	AAC	TGT	AGG	AAC

Reading direction  $\longrightarrow$

Document 2

<b>Codons</b>	UAA	UCU	ACA	AUU	GAU	CUU	UGU	UUU
	UAG	UCC	ACG	AUC	GAC	UUG	UGC	UUC
<b>Amino acids</b>	Stop	Ser	Thr	Ile	ac.Asp	Leu	Cys	Phe

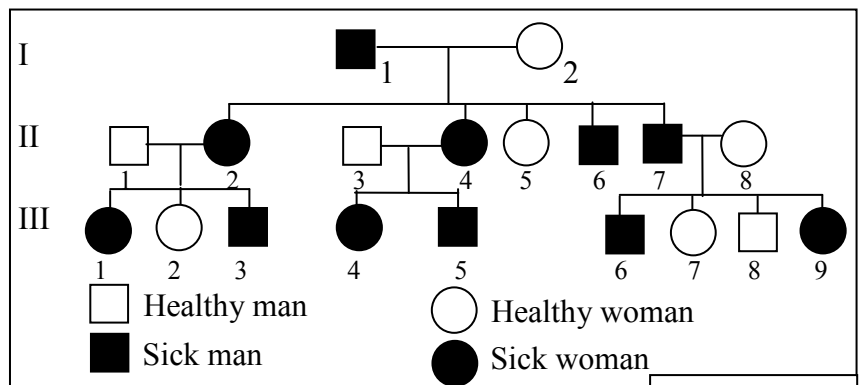
Document 3

2. Based on documents 2 and 3, **give** mRNA and the amino acids sequences corresponding to the normal allele and the abnormal allele, then **show** the genetic origin of neurofibromatosis type 1. (1.5 pts)

• Document 4 presents a pedigree of a family whose members are affected by the neurofibromatosis type 1.

3. Based on document 4 and knowing that  $I_2$  is homozygote:

a. **Show** that the allele responsible for the disease is dominant and autosomal (non-sexual chromosome). (0.5pts)



Document 4

b. Use Punnett Square to **determine** the probability that couple  $II_1$  and  $II_2$  would give birth to a healthy child (0.75pts)

(Use the symbols  $M$  and  $m$  for the two alleles of the gene referred to above)

• Neurofibromatosis type 1 is a widely spread disease. It affects one person out of 3500 individuals of a given population. Let's suppose that this population abides by the Hardy-Weinberg equilibrium:

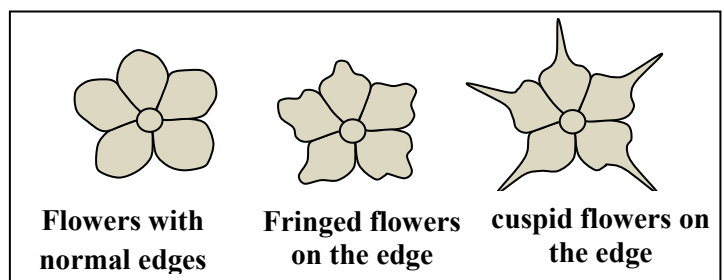
4. a. **Calculate** the frequency of the allele responsible for the disease and that of the normal allele. (0.5 pt)

b. **Calculate** the frequency of the heterozygote individuals. (0.5pts)

### Exercise 3 (3.25 pts)

Phloxes are herbaceous plants whose flowers show a wide variety of colors and shapes, a fact which makes them so important in horticulture.

❖ To study the transmission of two hereditary traits; the colors and shapes of flowers in phloxes, we suggest the following data : Phloxes have a variety of colors, including white and cream colors, and the edges of their petals can be of different shapes as shown in the document on the right hand side.



Flowers with normal edges

Fringed flowers on the edge

cuspid flowers on the edge

The following table presents results of crosses realized in phloxes.

Crosses	Cross I	Cross II
Parents P <sub>1</sub> X P <sub>2</sub>	between plants with white flower color and plants with cream flower color	between plants with normal flower shapes and plants with cuspid flower shapes
Generation F <sub>1</sub>	F <sub>1</sub> : plants with white flowers	F' <sub>1</sub> : plants with fringed flowers

1. What do you **deduce** from the results of the two crosses I and II? **(1 pt)**

**Cross III** - realized by the horticulturist between plants of pure lineage: plants with white flowers and regular shapes and plants with cream color flowers and cuspid shapes. This horticulturist wants to produce plants with cream color flowers and fringed shapes because the latter are easy to sell.

2. Bearing in mind that the two genes are independent,

a. Give the genotype of plants of generation F<sub>1</sub> (derived from Cross III), **(0,25 pts)**

b. Determine the theoretical results of generation F<sub>2</sub> derived from the cross between plants of generation F<sub>1</sub>. Justify your answer using Punnett Square **(1,25 pts)**

3. a. Give the genotype of plants that the horticulturist wants to produce **(0.25 pts)**

b. Based on the obtained genotypes of generation F<sub>2</sub>, suggest the cross that allows to obtain the greatest proportion of the desired phenotype. Justify your answer. **(0.5 pts)**

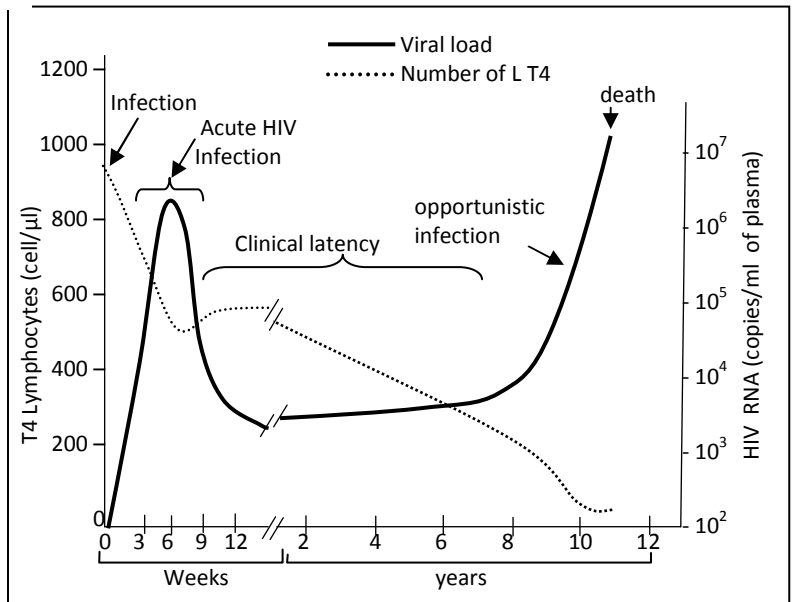
Use the following symbols: -*B* and *b* for alleles responsible for the white color of flowers ;  
-*C* and *c* for alleles responsible for the cuspid shape of flowers.  
-*N* and *n* for alleles responsible for the normal shape of flowers.

#### Exercise 4 (3.75 pts)

The infection by HIV (Human immunodeficiency virus) goes through many stages. The last stage is AIDS (acquired immunodeficiency syndrome) that is characterized by the appearance of opportunistic infections.

Knowledge of the immune system and the study of the infected individuals' reactions to HIV could help scientists to consider a vaccine against HIV. Document 1 shows the evolution of the number of lymphocytes T<sub>4</sub> and the virus load after infection by HIV.

**NB.** The viral load corresponds to the concentration of virus in blood and is reported as the number of viral RNA per millimeter of plasma.



Document 1

1. Based on document 1, describe the evolution of the number of lymphocytes T<sub>4</sub> and the viral load, then deduce the effect of HIV infection on the immune system. **(1.25pts)**

❖ The scientific community nowadays agree that for the vaccine to be effective against HIV, it must stimulate specific immune responses. To develop a vaccine, researchers have realized the following study: To test the vaccine, two batches of uninfected monkeys by HIV are used:

- The monkeys in the first batch received a series of five injections of vaccine;
- The monkeys of the second batch did not receive any injection.
- Afterwards, the two batches of monkeys were exposed to the virus.

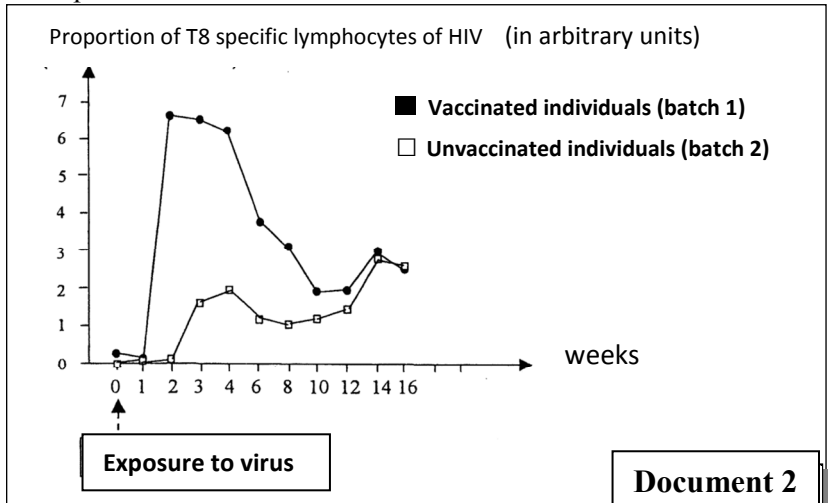
We evaluate the proportion of specific lymphocytes T<sub>8</sub> with regard to HIV in the monkeys' blood. Document 2 presents the results obtained.

2. Compare the evolution of proportions of specific lymphocytes T<sub>8</sub> with regard to HIV between the vaccinated monkeys and the unvaccinated monkeys, then deduce the characteristic of the immune response explaining the observed difference. (0.75 pts)

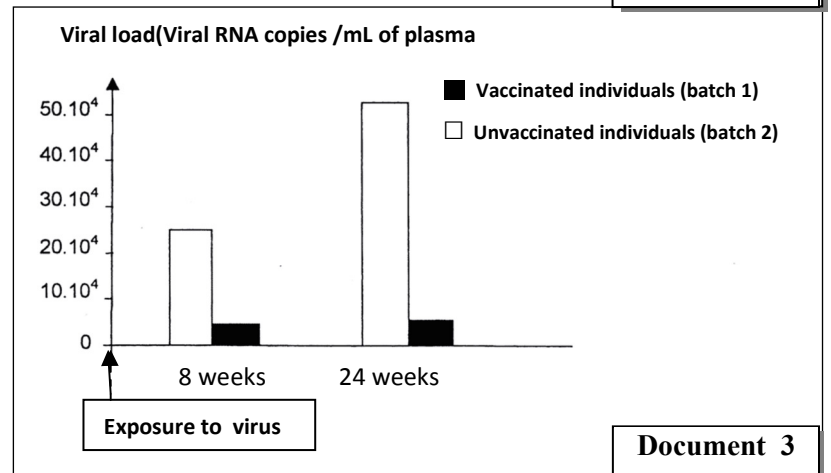
-We measured the viral load in monkeys of the two batches after 8 and 24 weeks of exposure to the virus. Document 3 presents the results of the measurements.

3. Compare the viral load in vaccinated and unvaccinated monkeys, then deduce the action of the experimental vaccine on the viral load. (0.75 pts)

❖ The study of the destruction mechanism of lymphocytes T<sub>4</sub> infected with HIV, and destroyed by lymphocytes cytotoxic T, allows to identify two mechanisms leading to the death of the target cell. Document 4 presents these two mechanisms.

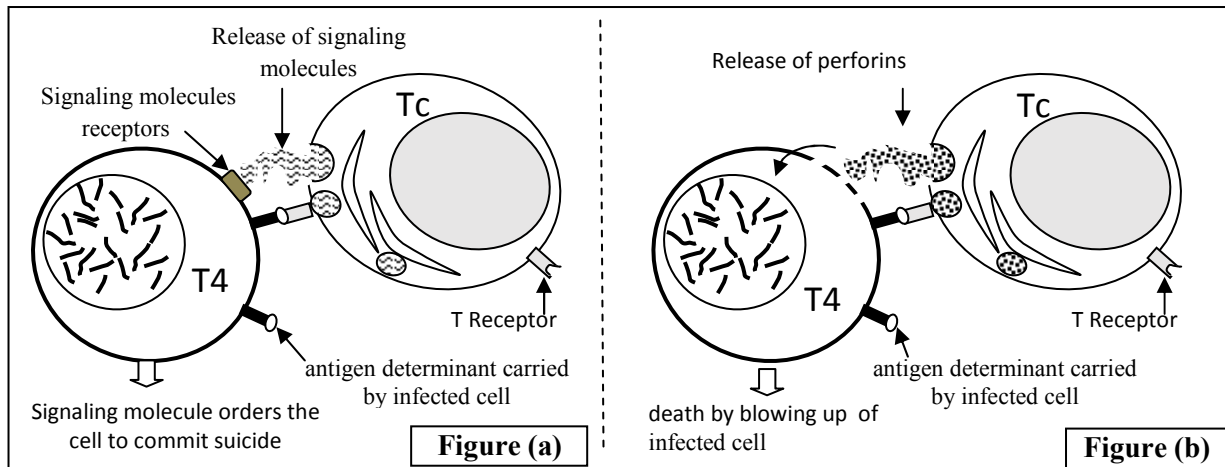


Document 2



Document 3

**NB. Despite the destruction of lymphocytes T<sub>4</sub> infected with HIV, the monkeys continued to produce other uninfected lymphocytes T<sub>4</sub>.**



Document 4

4. Based on document 4 and your answers to questions 2 and 3, explain the action mechanism of the tested vaccine on monkeys. (1 pt)









**Exercise 4 (3,75 pts)**

<b>1</b>	<p><b>Description of obtained results :</b>          The number of lymphocytes T4 gradually decreases after infection by HIV. It goes from 900 cells/<math>\mu</math>l to 50 cells/<math>\mu</math>l after 10 years of infection...          Viral load has rapidly increased after infection to reach (between <math>10^6</math> and <math>10^7</math> copies/ml of plasma) in the 6<sup>th</sup> week of infection. After that, it has decreased to stabilize at a value of (<math>10^3</math> and <math>10^4</math> copies/ml of plasma). After 8 years, it has again increased to reach a value upper to <math>10^7</math> copies/ml of plasma .....</p> <p><b>Deduction</b>          HIV infection <math>\rightarrow</math> decrease LT4 <math>\rightarrow</math> decrease of the immune defense system of individual organisms <math>\rightarrow</math> organism exposed to opportunistic diseases (weakening the immune system)</p>	0.5  0.5  0.25
<b>2</b>	<p>In vaccinated monkeys, unlike in unvaccinated ones, the production of LT8 is faster (after one week of exposure to HIV instead of two weeks for unvaccinated monkeys) and more intense (acute at 7 instead of 2 weeks for the unvaccinated monkeys)  <b>Deduction:</b> The characteristic is immunological memory .....</p>	0.5  0.25
<b>3</b>	<p><b>Comparison of the proportions of viral load in monkeys:</b>          In week 8 of virus exposure, the viral load in unvaccinated monkeys is 5 times higher than in vaccinated monkeys.          After week 24, the viral load has increased just a little in vaccinated monkeys while in the unvaccinated monkeys it has doubled.  <b>Deduction:</b> The vaccine inhibits the multiplication of HIV .....</p>	0.25  0.25  0.25
<b>4</b>	<p><b>Explantation :</b>          The use of the vaccine leads to the increase of LTc <math>\rightarrow</math> destruction of the lymphocyte LT4 infected by HIV in two ways: release of perforin and granzyme or/and the signals provoking apoptosis of the infected cell <math>\rightarrow</math> decrease of the number of infected lymphocytes LT4 <math>\rightarrow</math> decrease of viral load <math>\rightarrow</math> to avoid the appearance of opportunistic diseases.</p>	0.25  0.5  0.25